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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/632,054	07/31/2003	Robert E. Richard	02-465	9964
27774 MAYER & WII	7590 08/21/200 LLIAMS PC	EXAMINER		
251 NORTH A	·	HUGHES, ALICIA R		
2ND FLOOR WESTFIELD, I	NJ 07090		ART UNIT	PAPER NUMBER
			1614	
			MAIL DATE	DELIVERY MODE
			08/21/2009	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

## Advisory Action Before the Filing of an Appeal Brief

Application No.		Applicant(s)	
	10/632,054	RICHARD ET AL.	
	Examiner	Art Unit	
	ALICIA R. HUGHES	1614	

	ALICIA N. HOGHES	1014	
The MAILING DATE of this communication appe	ars on the cover sheet with the c	correspondence add	ress
THE REPLY FILED <u>07 July 2009</u> FAILS TO PLACE THIS APPL	ICATION IN CONDITION FOR AL	LOWANCE.	
1. The reply was filed after a final rejection, but prior to or on application, applicant must timely file one of the following application in condition for allowance; (2) a Notice of Appelor Continued Examination (RCE) in compliance with 37 C periods:	replies: (1) an amendment, affidavi eal (with appeal fee) in compliance	t, or other evidence, w with 37 CFR 41.31; or	hich places the (3) a Request
a) The period for reply expiresmonths from the mailing	date of the final rejection.		
b) The period for reply expires on: (1) the mailing date of this A no event, however, will the statutory period for reply expire la	dvisory Action, or (2) the date set forth ater than SIX MONTHS from the mailing	g date of the final rejection	n.
Examiner Note: If box 1 is checked, check either box (a) or (IMONTHS OF THE FINAL REJECTION. See MPEP 706.07(f	r).		
Extensions of time may be obtained under 37 CFR 1.136(a). The date of have been filed is the date for purposes of determining the period of extunder 37 CFR 1.17(a) is calculated from: (1) the expiration date of the s set forth in (b) above, if checked. Any reply received by the Office later may reduce any earned patent term adjustment. See 37 CFR 1.704(b). NOTICE OF APPEAL	ension and the corresponding amount of hortened statutory period for reply origi	of the fee. The appropria nally set in the final Offic	ate extension fee e action; or (2) as
2. ☐ The Notice of Appeal was filed on . A brief in comp	liance with 37 CEP 41 37 must be	filed within two month	e of the date of
filing the Notice of Appeal was filed off A brief in completing the Notice of Appeal (37 CFR 41.37(a)), or any exter Notice of Appeal has been filed, any reply must be filed with the state of the Notice of Appeal has been filed.	nsion thereof (37 CFR 41.37(e)), to	avoid dismissal of the	
<u>AMENDMENTS</u>			
<ol> <li>The proposed amendment(s) filed after a final rejection, to</li> <li>They raise new issues that would require further cor</li> </ol>	nsideration and/or search (see NO		cause
(b) They raise the issue of new matter (see NOTE below	•		
<ul><li>(c) ☐ They are not deemed to place the application in bett</li><li> appeal; and/or</li></ul>	er form for appeal by materially red	ducing or simplifying t	ne issues for
(d) ☐ They present additional claims without canceling a c	corresponding number of finally reje	ected claims.	
NOTE: (See 37 CFR 1.116 and 41.33(a)).			
4. 🔲 The amendments are not in compliance with 37 CFR 1.12		mpliant Amendment (	PTOL-324).
5. Applicant's reply has overcome the following rejection(s):			
<ol> <li>Newly proposed or amended claim(s) would be all non-allowable claim(s).</li> </ol>			_
7.  For purposes of appeal, the proposed amendment(s): a) how the new or amended claims would be rejected is proved the status of the claim(s) is (or will be) as follows: Claim(s) allowed:		l be entered and an e	xplanation of
Claim(s) objected to:			
Claim(s) rejected: <u>1,3-13,16-18 and 30</u> . Claim(s) withdrawn from consideration: <u>15 and 19-29</u> .			
AFFIDAVIT OR OTHER EVIDENCE			
<ol> <li>The affidavit or other evidence filed after a final action, but because applicant failed to provide a showing of good and was not earlier presented. See 37 CFR 1.116(e).</li> </ol>			
9. The affidavit or other evidence filed after the date of filing a entered because the affidavit or other evidence failed to or showing a good and sufficient reasons why it is necessary	vercome <u>all</u> rejections under appea	al and/or appellant fail	s to provide a
10. ☐ The affidavit or other evidence is entered. An explanation REQUEST FOR RECONSIDERATION/OTHER	n of the status of the claims after er	ntry is below or attach	ed.
11. The request for reconsideration has been considered but See Continuation Sheet.	does NOT place the application in	condition for allowan	ce because:
<ul><li>12. ☐ Note the attached Information <i>Disclosure Statement</i>(s). (</li><li>13. ☐ Other:</li></ul>	PTO/SB/08) Paper No(s)		
10. [_] Outer			
	/Raymond J Henley III/ Primary Examiner, Art U	nit 1614	

Continuation of 11. does NOT place the application in condition for allowance because:

With regard to the obviousness double patenting rejection over U.S. Patent No. 7,241,455 [the "Richard patent"], Applicant argues that unlike the Richard patent, the instant invention teaches the opposite of cross-linking to reduce molecular weight. This argument remains ubpersuasive, because both sets of claims substantially retain the same limitations, bringing the instant invention within the purview of the first invention.

With regard to the 103(3) rejection, the Applicants continue to argue that none of the references alone or in combination teach or suggest the current invention. Applicants have focused on a number of distinctions in individual references to support their claim that the instant invention is not prima facie obvious. The Examiner instead focuses on why the references may be read in tandem to reach a conclusion of obviousness by one of ordinary skill in the art.

Importantly, these 2.5 Mrads do read on the instant invention, which calls for a radiation dose that is at least 100,000 rads and claim 30, which has a range of 1 Mrad to 10 Mrads.

Further, Applicants' argue that there is no teaching or suggestion in Phan et al of polymers in which radiation is used to increase release of a therapeutic agent from a polymer is unfounded. Rather, as noted prior, it logically follows that the same therapeutic agent comprising the same polymeric release region is used in the same host, subject to a radiation dose that gives the same effect to the same patient population (Please see Col. 6, lines 29-33). And further as noted prior, by Applicants' own account, "when polymers are exposed to radiation, at least two reactions are believed to occur" and "[c]rosslinking generally results in ...[and c]hain scission, on the other hand generally results in ...While polymers may display both types of reactions, one type of reaction will typically dominate. For increased release, it is preferred to use polymers in which chain scission reactions dominate." (Emphasis added).

Applicants also argue that since Cruise et al teach hydrogels, that it is not germane to the instant invention. This one teaching does not negate the other teachings in Cruise et al as noted prior in the record and further, all other contentions with regard to Cruise et al and its teachings are but allegations that lack factual support. As a result, they are assigned no patentable weight.

Applicants also argue that there is no support to combine the teachings of Phan et al with the teachings of Pinchuk, because Pinchuk does not teach or suggest the use of any radiation treatment. By Applicants' own admission, however, Pinchuk does disclose the release of therapeutic agents over time. Given the state of the art, particularly the release of therapeutic agents over time in stent systems, and as well, the reasons made previously of record for combining the references, the Applicants' assertion that there is not support to modify the teachings of Phan et al with the teachings of Pinchuk is but an allegation lacking factual support.

With regard to the Furst reference, Applicants also argue that support for its use cannot be sustained, most notably because the amount of radiation used in Furst is less than 2000 rads. The Furst reference is not utilized as a single reference to anticipate the instant invention. Rather, it is used in concert with, to modify other references that teach stent technology. It would not be unreasonable or outside the motivation of one of ordinary skill in the art to modify Furst et al by the teachings in the other cited references to arrive at the radiation dosage taught by the instant invention.